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FLUID COMPOSITIONS CONTAINING REFRIGERATOR OILS AND CHLORINE-FREE FLUOROCARBON REFRIGERANTS

ADD 1-10-03 1010-1-3 1010-1-3 1011-1-3 This application is a Continuation-in-part of Ser. No. 08/539,001 filed October 4, 1995 which is a Continuation of Ser. No. 08/193,281 filed February 8, 1994 which is Continuation-in-part of Ser. No. 08/019,177 filed January 28, 1993 which is a Continuation of Ser. No. 07/634,054 filed December 26, 1990 now standard.

Background of the Invention

1. Field of the Invention

This invention relates to a lubricating oil for compressors of refrigerators using therein a hydrogen-containing halogenocarbon as a refrigerant (the oil being hereinafter referred to as "a refrigerator oil for use with a hydrogen-containing halogenocarbon refrigerant") and, more specifically, it relates to such a refrigerator oil which comprises a specific ester as a base oil and is superior in various properties.

2. Prior Art Background of The Prior Art

Generally, naphthenic mineral oils, paraffinic mineral oils, alkylbenzenes, polyglycolic oils, ester oils and mixtures thereof, which have each a kinematic viscosity of 10 - 200 cSt at 40°C, as well as these oils incorporated with suitable additives have been used as refrigerator oils.

On the other hand, chlorofluorocarbons (CFCS) type refrigerants, such as CFC-11, CFC-12, CFC-113 and HCFC-22, have been used for refrigerators.

Of these CFCS, CFCS such as CFC-11, CFC-12 and CFC-113, which are obtained by substituting all the hydrogen

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atoms of hydrocarbons thereof by halogen atoms including chlorine atoms, may lead to the destruction of the ozone layer, and therefore, the use of the CFCS has been controlled. Accordingly, hydrogen-containing

halogenocarbons, such as HFC-134a and HFC-152a, have been 5 being used as substitutes for CFCs. HFC-134a is especially promising as a substitute refrigerant since it is similar in thermodynamic properties to CFC-12 which has heretofore been used in many kinds of refrigerators of home cold-storage chests, air-conditioners and the like. 10

Refrigerator oils require various properties, among which their compatibility with refrigerants is extremely important in regard to lubricity and system efficiency in However, conventional refrigerator oils refrigerators. comprising, as the base oils, naphthenic oils, paraffinic oils, alkylbenzenes, heretofore known ester oils and the like, are hardly compatible with hydrogen-containing halogenocarbons such as HFC-134a. Therefore, if said conventional refrigerator oils are used in combination with HFC-134a, the resulting mixture will separate into two layers at normal temperature so as to degrade the oilreturnability which is the most important within the refrigeration system and cause various troubles such as a decrease in refrigeration efficiency, the deterioration of lubricity and the consequent seizure of the compressor within the system whereby the refrigerator oils are made

25 unsuitable for use as such. In addition, polyglycolic oils

are also known as refrigerator oils for their high viscosity index and are disclosed in, for example, JP-A-57-42119 and JP-A-61-52880 and JP-A-57-51795. However, the polyglycolic oils disclosed in these prior art publications are not fully compatible with HFC-134a thereby raising the same problems as above and rendering them unusable.

Further, US-A-4,755,316 discloses polyglycolic refrigerator oils which are compatible with HFC-134a and US-A-4,851,144 discloses refrigerator oils comprising a mixture of an ester and a polyglycol which are compatible with HFC-10 134a. In addition, the present inventors developed polyglycolic refrigerator oils which have excellent compatibility with HFC-134a as compared with conventional known refrigerator oils, filed an application for a patent for the thus developed polyglycolic refrigerator oils and 15 have already obtained a patent (US-A-4,948,525) therefor. It has been found, however, that the polyglycolic oils raise problems as to their high compatibility with water and inferior electrical insulating property.

20 On the other hand, refrigerator oils used in compressors of home refrigerators and the like are required to have a high electrical insulating property. Among the known refrigerator oils, alkylbenzenes and the mineral oils have the highest insulating property, but they are hardly compatible with hydrogen-containing halogenocarbons such as HFC-134a as mentioned above. WO 90/12849 describes a composition comprising a hydrogen-containing halogenocarbon

and a specific ester lubricant. No refrigerator oil having both high compatibility with hydrogen-containing halogenocarbons such as HFC-134a and a high insulating property has been developed prior to the present invention.

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Summary of the Invention

The present inventors made various intensive studies in attempts to develop refrigerator oils which can meet the aforesaid requirements and, as the result of their studies, they found that esters having specific structures have excellent compatibility with hydrogen-containing halogenocarbons such as HFC-134a, and a high electrical insulating property as well as excellent lubricity. This invention is based on this finding.

The object of this invention is to provide refrigerator oils for use with hydrogen-containing halogenocarbons refrigerants, the oils comprising as a major component (or a base oil) at least one kind of ester having a specific structure and an epoxy compound and having excellent compatibility with hydrogen-containing halogenocarbons such as HFC-134a, high electrical insulating property, high wear resistance, low hygroscopicity, and high thermal and chemical stability.

The present invention provides a refrigerator oil for use in compressors using therein a hydrogen-containing halogenocarbon as a refrigerant, consisting essentially of

as a base oil at least one ester selected from the group consisting of:

[I] a pentaerythritol ester represented by the general formula (1)

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wherein R¹-R⁴ may be identical with or different from each other and are each selected from the group consisting of straight-chain alkyl groups having 3 to 11 carbon atoms, branched-chain alkyl groups having 3 to 15 carbon atoms and cycloalkyl groups having 6 to 12 carbon atoms and a is an integer of 1 to 3;

[II] a polyol ester represented by the general formula (2)

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$$R^{6}C-O = CH_{2} C CH_{2}O = CR^{7}$$

$$CH_{2}O CH_{2}O CR^{7}$$

$$CR^{7}C CR^{7}C CR^{$$

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wherein R^s - R^7 may be identical with or different from each other and are each selected from the group consisting of straight-chain alkyl groups having 3 to 11 carbon atoms,

branched-chain alkyl groups having 3 to 15 carbon atoms and cycloalkyl groups having 6 to 12 carbon atoms, the straight-chain alkyl groups being present in a ratio of not more than 60 % of the total alkyl groups, R^8 is selected from the group consisting of methyl, ethyl and propyl groups, and b is an integer of 1 to 3;

[III] an ester represented by the general formula (3)

wherein X^1 is a group represented by the general formula $-0R^{1\,1}$ or $-0-R^{1\,2}-0-C-R^{1\,3}$, X^2 is a group represented 0

by the general formula $-C-R^{14}$ or $-C-(R^{15})_e$ $-C-OR^{16}$, R^9 and \parallel 0 0

R¹⁵ are each a divalent hydrocarbon group having 1 to 8 carbon atoms, R¹⁰ and R¹² are each a divalent saturated hydrocarbon group having 2 to 16 carbon atoms, R¹¹ and R¹⁶ are each a branched-chain alkyl group having 3 to 15 carbon atoms, R¹³ and R¹⁴ are each a branched-chain alkyl group

having 3 to 14 carbon atoms, c and e are each an integer of 0 or 1 and d is an integer of 0 to 30; and

[IV] a polyol ester obtained by the synthesis of, as raw materials, (a) a neopentyl type polyhydric alcohol having 5 to 6 carbon atoms and 3 to 4 hydroxyl groups, (b) a straight-chain monocarboxylic acid and/or a branched-chain monocarboxylic acid, the branched-chain monocarboxylic acid being present in a ratio of not less than 50 mol% of the total monocarboxylic acids, (c) a dicarboxylic acid: and further comprising at least one epoxy compound.

First, the pentaerythritol esters [I] will be 10 explained in detail. In the formula (1), R^1-R^4 may be identical with, or different from, each other and are each a substituent group selected from the group consisting of straight-chain alkyl groups having 3-11 carbon atoms, preferably 3-7 carbon atoms, branched-chain alkyl groups having 3-15 carbon atoms, preferably 4-11 carbon atoms and cycloalkyl groups having 6-15 12 carbon atoms, preferably 6-8 carbon atoms. cycloalkyl groups in this invention may include alkylcycloalkyl groups. In addition, the ratio of the straight-chain alkyl groups to the total alkyl groups is 20 preferably not more than 60%, more preferably not more than Further, a is an integer of 1-3. Thus, the formula(1) illustrates monopentaerythritol esters, dipentaerythritol

R¹-R⁴ are each exemplified by n-propyl group, n-butyl group, n-pentyl group, n-hexyl group, n-heptyl group, n-octyl group, n-nonyl group, n-decyl group, n-undecyl group,

esters and tripentaerythritol esters.

hexylcyclohexyl group.

iso-propyl group, iso-butyl group, iso-pentyl group, isohexyl group, iso-heptyl group, iso-octyl group, iso-nonyl group, iso-decyl group, iso-undecyl group, iso-dodecyl group, iso-tridecyl group, iso-tetradecyl group, isopentadecyl group, cyclohexyl group, cycloheptyl group, cyclooctyl group, cyclononyl group, cyclodecyl group, cycloundecyl group, cyclododecyl group, methylcyclohexyl group, ethylcyclohexyl group, propylcyclohexyl group, butylcyclohexyl group, pentylcyclohexyl group or

The pentaerythritol esters [I] are esters of pentaerythritol, dipentaerythritol or tripentaerythritol and a monocarboxylic acid, and are ordinarily obtained by the reaction of pentaerythritol, dipentaerythritol,

15 tripentaerythritol or a mixture thereof with a mixture of at least one carboxylic acid having the aforementioned alkyl group.

Secondly, the polyol esters [II] will be explained in In the formula (2), R^5-R^7 may be identical with, or different from, each other and are each a group selected from the group consisting of straight-chain alkyl groups having 3-11 carbon atoms, preferably 3-7 carbon atoms, branched-chain alkyl groups having 3-15 carbon atoms, preferably 4-11 carbon atoms and cycloalkyl groups having 6-25 12 carbon atoms, preferably 6-8 carbon atoms. cycloalkyl groups in this invention may include alkylcycloalkyl groups. With respect to the above R^5-R^7 ,

the ratio of the straight-chain alkyl groups to the total alkyl groups (including cycloalkyl groups) is not more than 60%, preferably not more than 50%. In addition, R⁸ is a group selected from the group consisting of methyl group, ethyl group and propyl group. Further, b is an integer of 1-3. Thus, the formula (2) illustrates trimethylolethane esters, trimethylolpropane esters, trimethylolbutane esters and dimers-trimers thereof.

 R^5-R^7 are each exemplified by n-propyl group, n-butyl group, n-pentyl group, n-hexyl group, n-heptyl group, n-10 octyl group, n-nonyl group, n-decyl group, n-undecyl group, iso-propyl group, iso-butyl group, iso-pentyl group, isohexyl group, iso-heptyl group, iso-octyl group, iso-nonyl group, iso-decyl group, iso-undecyl group, iso-dodecyl group, iso-tridecyl group, iso-tetradecyl group, iso-15 pentadecyl group, cyclohexyl group, cycloheptyl group, cyclooctyl group, cyclononyl group, cyclodecyl group, cycloundecyl group, cyclododecyl group, methylcyclohexyl group, ethylcyclohexyl group, propylcyclohexyl group, 20 butylcyclohexyl group, pentylcyclohexyl group or hexylcyclohexyl group.

The polyol esters [II] are esters of
trimethylolethane, trimethylolpropane, trimethylolbutane or
a dimer or trimer thereof and a monocarboxylic acid, and are
ordinarily obtained by the reaction of trimethylolethane,
trimethylolpropane, trimethylolbutane, a dimer or trimer

thereof or a mixture thereof with a mixture of at least one carboxylic acid having the aforementioned alkyl group.

Thirdly, the esters [III] will be explained in In the formula (3), X^1 is a group represented by detail. the general formula $-0R^{11}$ or and X^2 is a group represented by the general formula, . In addition, R^9 and R^{15} are each a ordivalent hydrocarbon group having 1-8 carbon atoms, preferably 1-6 carbon atoms, R^{10} and R^{12} are each a divalent saturated hydrocarbon group having 2-16 carbon atoms, preferably 2-9 carbon atoms, $R^{1\,1}$ and $R^{1\,6}$ are each an alkyl group having 1-15 carbon atoms, preferably 1-12 carbon atoms, and R^{13} and R^{14} are each an alkyl group having 1-14 carbon atoms, preferably 1-11 carbon atoms. Further, c and e are each an integer of 0 or 1 and d is an integer of 0-30, preferably 1-30.

R⁹ and R¹⁵ are each exemplified by a methylene group, ethylene group, propylene group, trimethylene group, butylene group, tetramethylene group, pentamethylene group,

- hexamethylene group, heptamethylene group, octamethylene group, phenylene group, other unsaturated hydrocarbon groups or the like. R^{10} and R^{12} are each illustrated by nonamethylene group, decamethylene group, undecamethylene group, dodecamethylene group, tridecamethylene group,
- tetradecamethylene group, pentadecamethylene group, hexadecamethylene group, cyclohexylene group or the like in addition to the above alkylene groups (except for methylene

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group). R¹³ and R¹⁴ are each concretely exemplified by straight- or branched-chain alkyl groups, such as methyl group, ethyl group, propyl group, butyl group, pentyl group, hexyl group, heptyl group, octyl group, nonyl group, decyl group, undecyl group, dodecyl group, tridecyl group or tetradecyl group. R¹¹ and R¹⁶ are each illustrated by a straight- or branched-chain alkyl group, such as a pentadecyl group, in addition to any one of the above alkyl groups.

- The method for producing the esters [III] is not particularly limited. For example, a mixture of the esters [III] can be produced by the esterification reaction of (i) a diol having none of ether linkages in the branched-chains thereof with (ii) a dicarboxylic acid and (iii) a
- 15 monocarboxylic acid and/or (iv) a monohydric alcohol. The esters [III] are also produced by the reaction of the diol (i) with the monocarboxylic acid (iii) or the reaction of the dicarboxylic acid (ii) with the monohydric alcohol (iii).
- The diols (i) used herein are those having 2-16 carbon atoms and are exemplified by alkylene glycols such as ethylene glycol, propylene glycol, butylene glycol, trimethylene glycol, tetramethylene glycol, pentamethylene glycol, hexamethylene glycol, heptamethylene glycol,
- octamethylene glycol, nonamethylene glycol, decamethylene glycol, neopentyl glycol, 2-ethyl-2-methyl-1,3-propanediol,

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2-methyl-2-propyl-1,3-propanediol, 2,2-dimethyl-1,3propanediol, 2-butyl-2-ethyl-1,3-propanediol, 3-methyl-1,5pentanediol, 2,4-pentanediol, 2-methyl-2,4-pentanediol, 2methyl-1,6-hexane, 1,4-cyclohexane dimethanol and 2,2-bis (4-hydroxycyclohexyl) propane.

The dicarboxylic acids (ii) used herein are those having 2-10 carbon atoms and are exemplified by saturated aliphatic dicarboxylic acids such as oxalic acid, malonic acid, succinic acid, glutaric acid, adipic acid, pimelic acid, suberic acid, azelaic acid, sebacic acid, 10 methylmalonic acid, ethylmalonic acid, dimethylmalonic acid, methylsuccinic acid, 2,2-dimethylsuccinic acid, 2,3dimethylsuccinic acid, 2-ethyl-2-methylsuccinic acid, 2methylglutaric acid, 3-methylglutaric acid, 3,3dimethylglutaric acid and 3-methyladipic acid; unsaturated aliphatic dicarboxylic acids such as maleic acid, fumaric acid, itaconic acid, citraconic acid and mesaconic acid; and

20 The monocarboxylic acids (iii) used herein are those having 2-15 carbon atoms and are exemplified by acetic acid, propionic acid, butanoic acid, pentanoic acid, hexanoic acid, heptanoic acid, octanoic acid, nonanoic acid, decanoic acid, undecanoic acid, dodecanoic acid, tridecanoic acid, 25

tetradecanoic acid, pentadecanoic acid, 3-methylbutanoic

aromatic dicarboxylic acids such as phthalic acid,

isophthalic acid and terephthalic acid.

acid, 2-methylbutanoic acid, 2-ethylhexanoic acid, 2,4-dimethylpentanoic acid, 3,3,5-trimethylhexanoic acid and benzoic acid.

The monohydric alcohols (iv) used herein are those having 1-15 carbon atoms and are exemplified by methanol, ethanol, propanol, butanol, pentanol, hexanol, heptanol, octanol, nonanol, decanol, unedecanol, dodecanol, tridecanol, tetradecanol, pentadecanol, isopropanol, isobutanol, 2-methyl-1-butanol, 2,2-dimethyl-1-propanol,

10 3,3-dimethyl-1-butanol, 2-methyl-1-pentanol, 3-methyl-1-pentanol, 2,4,4-trimethyl-1-pentanol, 2,2,4-trimethyl-1-pentanol, 2-ethyl-4-methyl-1-pentanol and 2-ethyl-1-hexanol.

The molecular weight of the ester [III] is not particularly limited, but the number average molecular weight of the ester [III] is in the range of preferably 200-3000, more preferably 300-2000, to improve the compressor in sealability

Fourthly, the polyol ester [IV] will be explained in detail.

The neopentyl type polyhydric alcohols (a) having 5-6 carbon atoms and 3-4 hydroxyl groups are illustrated by trimethylolethane, trimethylolpropane and pentaerythritol.

The monocarboxylic acids (b) preferably used herein are those having 2-15 carbon atoms and are exemplified by acetic acid, propionic acid, butanoic acid, pentanoic acid, hexanoic acid, heptanoic acid, octanoic acid, nonanoic acid,

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decanoic acid, undecanoic acid, dodecanoic acid, tridecanoic acid, tetradecanoic acid and pentadecanoic acid.

These carboxylic acids may be either straight-chain or branched-chain ones, but it is preferable to use the latter having a branched-chain structure in an amount of not less than 50 mol%, preferably not less than 60 mol%.

The dicarboxylic acids (c) used herein are those having 2-10 carbon atoms and are exemplified by saturated aliphatic dicarboxylic acids such as oxalic acid, malonic acid, succinic acid, glutaric acid, adipic acid, pimelic acid, suberic acid, azelaic acid, sebacic acid, methylmalonic acid, ethylmalonic acid, dimethylmalonic acid, methylsuccinic acid, 2,2-dimethylsuccinic acid, 2,3dimethylsuccinic acid, 2-ethyl-2-methylsuccinic acid, 2-**15** methylglutaric acid, 3-methylglutaric acid and 3methyladipic acid; unsaturated aliphatic dicarboxylic acids such as maleic acid, fumaric acid, itaconic acid, citraconic acid and mesaconic acid; and aromatic dicarboxylic acids such as phthalic acid, terephthalic acid and isophthalic 20 acid.

The esters [IV] are produced by the esterification reaction of the aforesaid component (a) with the component (b) and the component (c) under ordinary esterification conditions, for example, in the presence of an acid catalyst such as sulfuric acid and at an temperature of 100-180°C.

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The chemical structure of the ester [IV] according to the present invention is illustrated by the following general formula (6)

wherein X^3 and X^4 may be identical with, or different from, each other and are each an alkyl group having 1-4 carbon atoms or a group represented by the general formula $-CH_2-O-CR^{31}$; $R^{26}-R^{29}$ and R^{31} may be identical with, or different from each other and are each an alkyl group having 4-15 carbon atoms; R^{30} is an alkylene group having 1-8 carbon atoms; j is an integer of 0 or 1; and k is an integer of 1-5.

The molecular weight of the polyol ester [IV] is not particularly limited, but the number average molecular weight of the ester [IV] is in the range of preferably 200-3000, more preferably 300-2000, to improve the compressor in sealability.

The products obtained by the methods as mentioned above may be refined to remove the by-products and/or unreacted reactants, but the by-products and/or unreacted reactants may be present in small amounts in the

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refrigerator oils of the present invention as far as they do not impair the excellent performances thereof.

In the preparation of the refrigerator oils of the present invention, the esters [I]-[IV] mentioned above may be used singly, or jointly as a mixture of at least two kinds of the esters.

The kinematic viscosities of the esters according to the present invention are in the range of preferably 2-150 cSt, more preferably 5-100 cSt at 100° C.

The refrigerator oil of the present invention may comprise as the only base oil at least one member selected from the above esters [I]-[IV] and, as required, it may additionally comprise other base oils for refrigerator oils. Among the other base oils, preferable ones are illustrated as follows:

a polyoxyalkylene glycol or an ether thereof represented by the general formula (4)

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$$R^{17} - (OR^{19})_{f} - OR^{18}$$

wherein R¹⁷ and R¹⁸ are each a hydrogen atom or an alkyl group having 1-18 carbon atoms, R¹⁹ is an alkylene group having 2-4 carbon atoms and f is an integer of 5-70, and a polyoxyalkylene glycol glycerol ether represented by the general formula (5)

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$$H_{2}C-O+R \xrightarrow{23}O \xrightarrow{g} R$$

$$\downarrow \qquad \qquad \downarrow \qquad \qquad \downarrow$$

wherein $R^{20}-R^{22}$ are each a hydrogen atom or an alkyl group having 1-18 carbon atoms, $R^{23}-R^{25}$ are each an alkylene group having 2-4 carbon atoms and g-i are each an integer of 5-7.

These conventional oils represented by the formulae (4) and (5) may be used singly or jointly for adding to the refrigerator oil of this invention. Further, the refrigerator oil of this invention may be incorporated with paraffinic mineral oils, naphthenic mineral oils, poly α -olefins, alkylbenzenes and the like, but, in this case, the resulting mixed oil will be lowered in compatibility with hydrogen-containing halogenocarbons.

The amount of these conventional base oils so incorporated is not particularly limited as far as the excellent performances of the refrigerator oil of this invention are not impaired, but the esters [I]-[IV] should be present in the resulting mixed oil in a ratio of usually more than 50 % by weight, preferably not less than 70 % by weight of the total amount of the mixed oil.

To further improve the refrigerator oil of the present invention in thermal stability and chemical stability, it is incorporated with at least an epoxy

compound. The epoxy compound used herein is preferably selected from the group consisting of phenylglycidyl ether epoxy compounds, alkylglycidyl ether epoxy compounds, glycidyl ester epoxy compounds, aryloxirane compounds, alkyloxirane compounds, alicyclic epoxy compounds, epoxidized fatty acid monoesters and epoxidized vegetable oils.

The said phenylglycidyl ether epoxy compounds used herein include phenylglycidyl ether and derivatives thereof such as alkylphenylglycidyl ethers. The said alkylphenylglycidyl ethers are those having 1 to 3 alkyl groups having 1 to 13 carbon atoms, among which are preferred those having an alkyl group having 4 to 10 carbon atoms, such as n-butylphenylglycidyl ether, i-

- butylphenylglycidyl ether, sec-butylphenylglycidyl ether, tert-butylphenylglycidyl ether, pentylphenylglycidyl ether, hexylphenylglycidyl ether, heptylphenylglycidyl ether, octylphenylglycidyl ether, nonylphenylglycidyl ether and decylphenylglycidyl ether.
- The alkylglycidyl ether epoxy compounds include decylglycidyl ether, undecylglycidyl ether, dodecylglycidyl ether, tridecylglycidyl ether, tetradecylglycidyl ether, 2-ethylhexylglycidyl ether, neopentyl glycol diglycidyl ether, trimethylolpropane triglycidyl ether, pentaerythritol tetraglycidyl ether, 1,6-hexanediol diglycidyl ether, sorbitol polyglycidyl ether, polyalkylene glycol

monoglycidyl ethers and polyalkylene glycol diglycidyl ethers.

The glycidyl ester epoxy compounds include phenylglycidyl esters, alkylglycidyl esters and alkenylglycidyl esters with glycidyl 2,2-dimethyl octanoate, glycidyl benzoate, glycidyl acrylate, glycidyl methacrylate and the like being preferred.

The aryloxirane compounds include 1,2-epoxystyrene and alkyl-1,2-epoxystyrene.

The alkyloxirane compounds include 1,2-epoxybutane, 1,2-epoxypentane, 1,2-epoxyhexane, 1,2-epoxyheptane, 1,2-epoxyhexane, 1,2-epoxydecane, 1,2-epoxydecane, 1,2-epoxydecane, 1,2-epoxydecane, 1,2-epoxytridecane, 1,2-epoxytridecane, 1,2-epoxyhexadecane, 1,2-epoxyhexadecane, 1,2-epoxyhexadecane, 1,2-epoxyhexadecane, 2-epoxyhexadecane, 1,1,2-epoxyoctadecane, 2-

1,2-epoxyheptadecane, 1,1,2-epoxyoctadecane, 2-epoxynonadecane and 1,2-epoxyeicosane.

The alicyclic epoxy compounds include 1,2epoxycyclohexane, 1,2-epoxycyclopentane, 3,4epoxycyclohexylmethyl-3,4-epoxycyclohexane carboxylate, bis

20 (3,4-epoxycyclohexylmethyl) adipate, exo-2,3epoxynorbornane, bis (3,4-epoxy-6-methylcyclohexylmethyl)
adipate, 2-(7-oxabicyclo[4.1.0]hept-3-yl)-spiro(1,3-dioxane5,3'-[7]oxabicyclo[4.1.0]heptane, 4-(1'-methylepoxyethyl)1,2-epoxy-2-methylcyclohexane and 4-epoxyethyl-1,2epoxycyclohexane.

The epoxidized fatty acid monoesters include esters of an epoxidized fatty acid having 12 to 20 carbon atoms and

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an alcohol having 1 to 8 carbon atoms, phenol or an alkylphenol. In particular, butyl, hexyl, benzyl, cyclohexyl, methoxyethyl, octyl, phenyl or butylphenyl esters of epoxidized stearic acid may preferably be used.

The epoxidized vegetable oils include epoxidized compounds of vegetable oils such as soybean oil, linseed oil and cottonseed oil.

Among these epoxy compounds, the preferable ones include phenylglycidyl ether epoxy compounds, alkylglycidyl ether type epoxy compounds, glycidyl ester type epoxy compounds and alicyclic epoxy compounds.

It is desirable that these epoxy compounds be incorporated in the refrigerator oil of the present invention in a ratio of 0.1-5.0% by weight, preferably 0.2-2.0% by weight, of the total amount of the refrigerator oil.

The refrigerator oil composition according to this invention may be incorporated further with at least one kind of a phosphorus compound selected from the group consisting of phosphoric esters, acid phosphoric esters, amine salts of acid phosphoric esters, chlorinated phosphoric esters and phosphorous esters, to improve the oil composition in wear resistance and load resistance. These phosphorus compounds are esters of phosphoric acid or phosphorous acid and an alkanol or a polyether type alcohol, or derivatives of the esters. The phosphoric esters are exemplified by tributyl phosphate, triphenyl phosphate and tricresyl phosphate. The acid phosphoric esters are exemplified by ditetradecyl acid

phosphate, dipentadecyl acid phosphate, dihexadecyl acid phosphate, diheptadecyl acid phosphate and dioctadecyl acid phosphate. The amine salts of acid phosphoric esters are exemplified by salts of the above acid phosphoric esters and amines such as methylamine, ethylamine, propylamine, butylamine, pentylamine, hexylamine, heptylamine, octylamine, dimethylamine, diethylamine, dipropylamine, dibutylamine, dipentylamine, dihexylamine, diheptylamine, dioctylamine, trimethylamine, triethylamine, tripropylamine,

- and trioctylamine. The chlorinated phosphoric esters are exemplified by tris-dichloropropyl phosphate, tris-chloroethyl phosphate, polyoxyalkylene-bis[di(chloroalkyl)] phosphate and
- tris chlorophenyl phosphate. The phosphorous esters are exemplified by dibutyl phosphite, tributyl phosphite, dipentyl phosphite, tripentyl phosphite, dihexyl phosphite, trihexyl phosphite, diheptyl phosphite, triheptyl phosphite, dioctyl phosphite, trioctyl phosphite, dinonyl phosphite,
- didecyl phosphite, diundecyl phosphite, triundecyl phosphite, didodecyl phosphite, tridodecyl phosphite, diphenyl phosphite, triphenyl phosphite, dicresyl phosphite, tricresyl phosphite and mixtures thereof. These phosphorus compounds may be added to the refrigerator oil in a ratio of 0.1-5.0 % by weight, preferably 0.2-2.0 % by weight, of the

total amount of the refrigerator oil.

Of course, both of the aforementioned phosphorus compounds and epoxy compounds may be used jointly.

To further enhance the refrigerator oil of this invention in performances, the refrigerator oil may be incorporated, as required, with heretofore known additives for a refrigerator oil, which include phenol-type antioxidants such as di-tert.-butyl-p-cresol and bisphenol A; amine-type antioxidants such as phenyl-α-naphthylamine and N.N-di(2-naphthyl)-p-phenylenediamine; wear resistant 10 additives such as zinc dithiophosphate; extreme pressure agents such as chlorinated paraffin and sulfur compounds; oiliness improvers such as fatty acids; antifoaming agents such as silicone-type ones; and metal inactivators such as benzotriazole. These additives may be used singly or The total amount of these additives added is **15** iointly. ordinarily not more than 10 % by weight, preferably not more than 5 % by weight, of the total amount of the refrigerator oil.

20 least one of the esters [I]-[IV] as the base oil should have such viscosity and pour point as those which are normally suitable for an ordinary refrigerator oil, but they should desirably have a pour point of not higher than -10°C, preferably -20°C to -80°C, to prevent them from solidification at a low temperature. Further, they should desirably have a kinematic viscosity of not less than 2cSt, preferably not less than 3cSt at 100°C, to keep the

sealability of the compressor when used, while they should desirably have a kinematic viscosity of not more than 150cSt, preferably not more than 100cSt at 100°C, in view of their fluidity at a low temperature and the efficiency of heat exchange in the evaporator when used.

The refrigerants which may be used in refrigerators in which the lubricating oils (refrigerator oils) of the present invention are suitably used, include hydrogencontaining halogenocarbons such as alkane fluorides having 1-3 carbon atoms, preferably 1-2 carbon atoms and/or alkane 10 chloride fluoride having 1-3 carbon atoms, preferably 1-2 The said hydrogen-containing halogenocarbons carbon atoms. are exemplified by HFCs (chlorine-free type halogenocarbons) such as difluoromethane (HFC-32), trifluoromethane (HFC-23), **15** pentafluoroethane (HFC-125), 1,1,2,2-tetrafluoroethane (HFC-134), 1,1,1,2-tetrafluoroethane (HFC-134a), 1,1,1trifluoroethane (HFC-143a) and 1,1-difluoroethane (HFC-152a); HCFCs (chlorine-containing type halogenocarbons) such as monochlorodifluoromethane (HCFC-22), 1-chloro-1,1difluoroethane (HCFC-142b), dichlorotrifluoroethane (HCFC-123) and monochlorotetrafluoroethane (HCFC-124); and mixtures thereof. Among these hydrogen-containing halogenocarbons, the chlorine-free type halogenocarbons such as HFC-32, HFC-23, HFC-125, HFC-134, HFC-134a and HFC-152a,

25 are preferable in view of the environmental problems. The refrigerant used may suitably be selected from these halogenocarbons mentioned above depending on the purpose for which the resulting refrigerant is used as well as the properties which are desirable for the resulting refrigerant. The preferable refrigerants are exemplified by HFC-134a; a mixture of HFC-134a (60-80wt%) and HFC-32 (40-20wt%); a mixture of HFC-32 (50-70wt%) and HFC-125 (50-30wt%); a mixture of HFC-134a (60wt%), HFC-32 (30wt%) and HFC-125 (10wt%); a mixture of HFC-134a (52wt%), HFC-32 (23wt%) and HFC-125 (25wt%); and a mixture of HFC-143a (52wt%), HFC-125 (44wt%) and HFC-134a (4wt%).

When the refrigerator oil of the present invention is used in a refrigerator, it is usually present in the form of a fluid composition for the refrigerator, which is a mixture of the refrigerator oil and a chlorine-free type halogenocarbon such as an alkane fluoride and/or an alkane chloride fluoride as mentioned above.

The present invention also relates to a fluid composition for a refrigerator, which comprises a chlorine-free type halogenocarbon refrigerant and a refrigerator oil according to the present invention.

The mixing ratio of the refrigerator oil and the refrigerant in the resulting composition is not particularly limited, but the refrigerator oil is usually comprised in an amount of 1-500 parts by weight, preferably in an amount of 2-400 parts by weight, based on 100 parts by weight of the refrigerant.

The refrigerator oils of the present invention are very excellent in compatibility with the hydrogen-containing

halogenocarbons as compared with the heretofore known refrigerator oils. Further, the refrigerator oils of the present invention are excellent because they have not only high compatibility with the hydrogen-containing

halogenocarbons and high electrical insulating property but also high lubricity, low hygroscopicity and high thermal and chemical stability.

The refrigerator oils of the present invention may particularly preferably be used in refrigerators, air
conditioners, dehumidifiers, cold-storage chests, freezers, freeze and refrigeration warehouses, automatic vending machines, showcases, cooling units in chemical plants, and the like which have a reciprocating or rotary compressor. Further, the above refrigerator oils may also preferably be used in refrigerators having a centrifugal compressor.

This invention will be better understood by the following Examples and Comparative Examples.

Examples 1-69 and Comparative Examples 1-70

First, base oils according to this invention (Base oils 1-23 and 30) and comparative base oils of conventional types (Base oils 24-29 and 31-49), which are used in the Examples and Comparative Examples, respectively, are illustrated as follows:

25

[Base oil 1] A tetraester of pentaerythritol (1 mol) and 2-ethylhexanoic acid (4 mol):

 $C \left\{\begin{array}{ll} CH_2-OOC-R \end{array}\right]_{\textbf{4}} \quad R \ ; \quad -CH-CH_2-CH_2-CH_2-CH_3 \\ C_2H_5 \end{array}$

[Base oil 2] A tetraester of pentaerythritol (1 mol) and 5 3,5,5-trimethylhexanoic acid (4 mol):

$$C = \begin{bmatrix} CH_{3} & CH_{3} \\ CH_{2} - OOC - R \end{bmatrix}_{4} \quad R \; ; \; -CH_{2} - CH_{2} - CH_{2} - CH_{3} \\ CH_{3} & CH_{3} \end{bmatrix}$$

10

[Base oil 3] A tetraester of pentaerythritol (1 mol), 2-ethylhexanoic acid (2 mol) and 3,5,5-trimethylhexanoic acid (2 mol)

15

[Base oil 4] A hexaester of dipentaerythritol (1 mol), n-hexanoic acid (3 mol) and 2,4-dimethylpentanoic acid (3 mol):

R; -CH2-CH2-CH2-CH2-CH3

20
$$\left\{R-COO-CH_{2}\right\}_{J_{3}}C-CH_{2}-O-CH_{2}-C\left\{CH_{2}-O-CO-R\right\}_{3}$$
 or $-CH-CH_{2}-CH-CH_{3}$ CH_{3} CH_{3}

[Base oil 5] A hexaester of dipentaerythritol (1 mol) and 3,5,5-trimethylhexanoic acid (6 mol):

R: $\left[R-COO-CH_{2}\right]_{3}C-CH_{2}-O-CH_{2}-C\left\{CH_{2}-O-CO-R\right]_{3}$ $-CH_{2}-CH-CH_{2}-C-CH_{3}$

5 [Base oil 6] A mixture of 50 parts by weight of the same ester as in Base oil 1 and 50 parts by weight of the same ester as in Base oil 5

[Base oil 7] A mixture of 30 parts by weight of the same 10 ester as in Base oil 2, 40 parts by weight of the same ester as in Base oil 5 and 30 parts by weight of the following octaester of tripentaerythritol (1 mol), 3-methylbutanoic acid (4 mol) and 3-methylpentanoic acid (4 mol):

15 $\left[R - COO - CH_2 \right]_3 C - CH_2 - O - CH_2 - \frac{1}{C} - CH_2 - O - CH_2 - C \left[CH_2 - O - CO - R \right]_3$ $CH_2 - O - CO - R$

R;
$$-CH_2 - CH - CH_3$$
 or $-CH_2 - CH - CH_2 - CH_3$ CH_3

[Base oil 8] A triester of trimethylolpropane (1 mol) and 2-ethylhexanoic acid (3 mol):

 $C_2H_5 - C \left\{ \begin{array}{l} CH_2 - OOC - R \\ \\ C_2H_5 \end{array} \right\}_3 \ R \ ; \ - CH - CH_2 - CH_2 - CH_2 - CH_3 \\ C_2H_5 \end{array}$

25

20

| |

[Base oil 9] A triester of trimethylolpropane (1 mol) and 3,5,5-trimethylhexanoic acid (3 mol):

$$\begin{array}{c} CH_{3} & CH_{3} \\ C_{2}H_{5}-C \left\{CH_{2}-OOC-R\right\}_{3} & R; -CH_{2}-CH-CH_{2}-CC-CH_{3} \\ CH_{3} & CH_{3} \end{array}$$

[Base oil 10] A triester of trimethylolpropane (1 mol), 2-10 ethylhexanoic acid (1.5 mol) and 3,5,5-trimethylhexanoic acid (1.5 mol)

[Base oil 11] A tetraester of di-(trimethylolpropane) (1 mol), n-hexanoic acid (2 mol) and 2,4-dimethylpentanoic acid (2 mol):

[Base oil 12] A tetraester of di-(trimethylolpropane) (1 mol) and 3,5,5-trimethylhexanoic acid (4 mol):

[Base oil 13] A mixture of 50 parts by weight of the same ester as in Base oil 8 and 50 parts by weight of the same ester as in Base oil 12

5 [Base oil 14] A mixture of 30 parts by weight of the same ester as in Base oil 9, 40 parts by weight of the same ester as in Base oil 12 and 30 parts by weight of the following octaester of tripentaerythritol (1 mol), 3-methylbutanoic acid (4 mol) and 3-methylpentanoic acid (4 mol):

10

$$\begin{bmatrix} CH_{2} - O - CO - R \\ R - COO - CH_{2} \end{bmatrix}_{3} C - CH_{2} - O - CH_{2} - \begin{bmatrix} CH_{2} - O - CH_{2} - C \\ CH_{2} - O - CO - R \end{bmatrix}_{3}$$

15

[Base oil 15] A diester of adipic acid and 2-ethyl-1-hexanol:

20

[Base oil 16] A diester of 3-methyl-1,5-pentanediol and 3,5,5-trimethylhexanoic acid:

25

i :

[Base oil 17] A complex ester having an average molecular weight of about 500 of 3-methyl-1,5-pentanediol, adipic acid and 3,5,5-trimethylhexanoic acid:

and

$$R_{2}-COO-C_{2}H_{4}-CH-C_{2}H_{4}-O = CO-(CH_{2})_{4}-COO-C_{2}H_{4}-CH-C_{2}H_{4}-O = CO-R_{2}$$

$$CH_{3}$$

$$CH_{3}$$

$$CH_{3}$$

$$CH_{3}$$

$$R \ge 1$$

 R^2 ; the same as in Base oil 16

20

[Base oil 18] A complex ester having the same composition as that in Base oil 17 except for having an average molecular weight of about 700

25

[Base oil 19] A complex ester having an average molecular weight of about 520 of 3-methyl-1,5-pentanediol, adipic acid and 2-ethyl-1-hexanol:

R1-0-CO-(CH2)4-COO-R1

and

$$R_1 - O \left\{ CO - (CH_2)_4 - COO - C_2H_4 - CH - C_2H_4 - O \right\} - CO - (CH_2)_4 - COO - R_1$$
 CH_3
 $n \ge 1$

5

[Base oil 20] A complex ester having the same composition as that in Base oil 19 except for having an average molecular weight of about 750

10

[Base oil 21] A mixture of the same complex ester as in Base oil 17 and 1.0 wt.% of a phosphoric ester type wear inhibitor

15 [Base oil 22] An ester having an average molecular weight of 690 and a kinematic viscosity at 100°C of 9.6 cSt, obtained by the reaction of 2 moles of trimethylolpropane with 1 mole of succinic acid, 1 mole of glutaric acid, 2 moles of iso-pentanoic acid and 2 moles of iso-hexanoic acid

20

25

[Base oil 23] An ester having an average molecular weight of 880 and a kinematic viscosity at 100 °C of 12.7 cSt, obtained by the reaction of 2 moles of pentaerythritol with 1 mole of succinic acid, 1 mole of glutaric acid, 3 moles of iso-pentanoic acid and 3 moles of iso-hexanoic acid

[Base oil 24] A naphthenic mineral oil

[Base oil 25] A branched-chain type alkylbenzene (average molecular weight: about 300)

[Base oil 26] Polyoxypropylene glycol monobutyl ether

(average molecular weight: about 500)

[Base oil 27] Polyoxypropylene glycol monobutyl ether (average molecular weight: about 1000)

10 [Base oil 28] Polyoxypropylene glycol (average molecular weight: about 700)

[Base oil 29] Polyoxypropylene glycol (average molecular weight: about 2000)

15

[Base oil 30] A tetraester of pentaerythritol (1 mol) and n-nonanoic acid (4 mol)

[Base oil 31] A tetraester of pentaerythritol (1 mol) and coconut oil

[Base oil 32] A triester of trimethylolpropane (1 mol) and n-nonanoic acid (3 mol)

25 [Base oil 33] A triester of trimethylolpropane (1 mol) and coconut oil

[Base oil 34] Polyoxyethyleneoxypropylene glycol mono n-butyl ether (number-average molecular weight = 670, kinematic viscosity at $100~^{\circ}\text{C} = 7.1~\text{mm}^2/\text{s}$, weight ratio of oxyethylene groups to oxypropylene groups in polyoxyalkylene chain = 1 : 1)

[Base oil 35] Propylene glycol derivative of polyoxy-propylene glycol mono methyl ether (number-average molecular weight = 1100, kinematic viscosity at 100 $^{\circ}$ C = 11.3 mm²/s):

$$\begin{array}{c}
\text{CH}_{2}^{-0-(C_{3}^{H}_{6}^{0})_{n}^{-CH}_{3}} \\
\text{CH}_{0}^{-0-(C_{3}^{H}_{6}^{0})_{n}^{-CH}_{3}} \\
\text{CH}_{0}^{-0-(C_{3}^{H}_{6}^{0})_{n}^{-CH}_{3}}
\end{array}$$

[Base oil 36] Glycerol derivative of polyoxypropylene glycol

15 mono methyl ether (number-average molecular weight = 1300,

kinematic viscosity at 100 °C = 12.8 mm²/s):

$$\begin{array}{c} ^{\rm CH}2^{-O-(C_3H_6O)}n^{-CH_3} \\ | \\ ^{\rm CH-O-(C_3H_6O)}n^{-CH_3} \\ | \\ ^{\rm CH_2-O-(C_3H_6O)}n^{-CH_3} \end{array}$$

20

5

[Base oil 37] Mixture of base oil 1 and base oil 27 (weight ratio of base oil 1 to base oil 27 = 70 : 30)

[Base oil 38] Mixture of base oil 1 and base oil 27 (weight ratio of base oil 1 to base oil 27 = 50 : 50)

[Base oil 39] Mixture of base oil 1 and base oil 34 (weight ratio of base oil 1 to base oil 34 = 30 : 70)

[Base oil 40] Mixture of base oil 1 and base oil 36 (weight ratio of base oil 1 to base oil 36 = 70 : 30)

[Base oil 41] Mixture of base oil 2 and base oil 36 (weight ratio of base oil 2 to base oil 36 = 50 : 50)

10 [Base oil 42] Mixture of base oil 3 and base oil 36 (weight ratio of base oil 3 to base oil 36 = 30 : 70)

[Base oil 43] Mixture of base oil 4 and base oil 27 (weight ratio of base oil 4 to base oil 27 = 50 : 50)

15

[Base oil 44] Mixture of base oil 5 and base oil 27 (weight ratio of base oil 5 to base oil 27 = 50 : 50)

[Base oil 45] Mixture of base oil 6 and base oil 27 (weight 20 ratio of base oil 6 to base oil 27 = 70 : 30)

[Base oil 46] Mixture of base oil 7 and base oil 27 (weight ratio of base oil 7 to base oil 27 = 30 : 70)

25 [Base oil 47] Mixture of base oil 1 and base oil 27 (weight ratio of base oil 1 to base oil 27 = 95 : 5)

[Base oil 48] Mixture of base oil 1 and base oil 34 (weight ratio of base oil 1 to base oil 34 = 95 : 5)

[Base oil 49] Mixture of base oil 30 and base oil 27 5 (weight ratio of base oil 30 to base oil 27 = 95 : 5)

the present invention each of which has a composition indicated in the following Table 1 were prepared and then evaluated for their performances that are their miscibility with HFC-134a, insulating property, wear resistance, hygroscopicity and thermal and chemical stability by the following respective test methods. The results thus obtained are indicated in Table 1.

15 For comparison, the mineral oil (Comparative Example 1), the alkylbenzene (Comparative Example 2), the polypropylene glycol monoalkyl ethers and derivatives thereof (Comparative Examples 3, 4, 34-45, 62 and 63), the polypropylene glycols (Comparative Examples 5 and 6), the 20 straight-chain fatty acid esters of polyol alcohols (Comparative Examples 8-10) which are free from epoxy compounds and have heretofore been used as refrigerator oils, mixed oils of the polypropylene glycol monoalkyl ethers or derivatives thereof with the esters of the present invention (Comparative Examples 46-61 and 64-70), mixed oils 25 of the tetraester of pentaerythritol, as well as Base oils 1-23 (Comparative Examples 11-33) which are free from epoxy

15

compounds, were evaluated for their performances in the same manner as in Examples 1-72. The results thus obtained are indicated in Table 2.

5 (Miscibility with HFC-134a)

0.2 g of the test oil of each of the Examples

(Examples 1-23 and 67-69) and the Comparative Examples

(Comparative Examples 1-33 and 70) and 1.8 g of the

refrigerant (HFC-134a) as well as 0.06 g of the test oil of

each of the Examples (Examples 1-23 and 67-69) and the

Comparative Examples (Comparative Examples 11-33 and 70) and

1.94 g of the refrigerant (HFC-134a) were sealed in a glass

tube having an inner diameter of 6 mm and a length of 220

mm. This glass tube was then placed in a thermostat

maintained at a predetermined low temperature or high

temperature to observe whether the refrigerant and the test

oil were miscible with each other, separated from each other

or made white-turbid.

20 (Insulating property)

The test oils were each measured for specific volume resistivity at 25°C in accordance with JIS C 2101.

(FALEX wear test)

The test oils were each applied to a test journal for measuring the amount of the test journal worn by having the journal run in at a test oil temperature of 100 °C under a

load of 150 lb for 1 minute and then running it under a load of 250 lb for 2 hours in accordance with ASTM D 2670.

(Hygroscopicity)

Thirty grams (30 g) of each of the test oils were placed in a 300-ml beaker, allowed to stand for 7 days in an air-conditioned bath maintained at a temperature of 30 °C and a humidity of 60 % and then measured for water content by the Karl-Fischer method.

10

(Thermal and chemical stability)

Thermal and chemical stabilities of the test oils were evaluated by the following sealed tube test with HFC-134a in accordance with JIS K 2211 4.9.

the refrigerant (HFC-134a) was sealed, together with copper, iron and aluminum catalysts, in a glass tube and then heated to 250°C for 168 hours to observe discoloration of the test oil and surface states of the catalysts and determine the degree of discoloration. The degree of discoloration of the test oil was classified into 9 grades ranging from 0 (no discoloration) to 8 (black-brown color).

Fifty grams (50 g) of each of the test oils (Examples 24-66 and Comparative Examples 34-69) indicated in the Tables were placed in a 100-ml beaker, and then allowed to stand for 168 hours in an air-conditioned bath maintained at a temperature of 25 °C and a relative humidity of 80%.

Next, an equi-volume mixture of each of these test oils and the refrigerant (HFC-134a) was sealed, together with copper, iron and aluminum catalysts, in a glass tube and then heated to $175~^{\circ}\text{C}$ for 168 hours to observe whether the state of the test oil has been changed or not.

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Table 1

	a)	 		Γ			-,																				بو	بو	بو.
	Appearance	of test oil				1		-		!	-	-	!	!				;		-	!	-		-	1	1	No change	No change	No change
			Al	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	:	1	:				
Sealed tube test	Catalyst		Fe	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	-	-	1				
			ß	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	1						
	0i1	color		1	_	-	-	1	-	-	-	-	, ,		1	1	1	1	1	-		1		0	-	_			-
Hygro- sconicity	30°C, 60%	€		0. 19	0.20	0.18	0.18	0.16	0.16	0.12	0.21	0.20	0.22	0.17	0.14	0.15	0.12	0.22	0.21	0.16	0.11	0.17	0.11	0.16	0.20	0.12	0.19	0.20	0.18
Falex test	journal	worn (mg)		26	25	92	21	19	21	19	27	27	25	23	20	52	20	23	20	16	11	=======================================	10	ç	19	16	!	;	-
Resisti-	925°C	(Dcm)		4.2×1014	3.8×1014	3.9×1014	2.9×1014	3.0×10^{14}	3.1×10 ¹⁴	2. 4×10 ¹⁴	4.5×10 ¹⁴	4.0×1014	4.1×10 ¹⁴	2.7×10¹⁴	2.8×10 ¹⁴	3.1×10 ¹⁴	2.7×1014	4.0×1014	3.9×10 ¹⁴	2.4×10 ¹⁴	1.4×10¹4	2.4×1014	1.1×1014	2.5×1014	2. 9×10^{14}	1.4×10¹4	4.2×1014	3.8×10 ¹⁴	3. 9×1014
mp. Range	(C)	Amount of	HFC-134a (97 wt%)	-45~CT**	-60~CT**	-45~CT	<-70~CT*	-18~30	-36~CT.	<-70~CT.	<-70~CT.	<-70~CT	<-70~CT**	-25~CT**	-38~17	-21~CT	-19~CT•	-38~CT**	-54~CT.	<-70~CT.	.L3~09-	<-70~CT**	-56~CT**	-70~CT•	<-50~CT.	<-50~CT**	-	1	-
Miscibile Temp. Range with HFC-134a (か)	1 0 11 1151	Amount of	HFC-134a (90 wt%)	-27~CT**	-32~CT.	-28~CT**	<-70~CT**	-7~75	-25~30	-5~86	-35~CT**	-45~CT**	-37~CT**	<-70~CT**	-11~75	-20~0	-7~88	-24~CT.	-30~CI.	-55~CT**	-34~-95	-47~CT**	-32~32	-55~CT	<-35~76	<-30~98		1	-
Kinematic	0100°C	(mm ² /s)		6.2	11.5	8.1	10.0	28.2	11.8	32. 3	3.7	9.4	7.1	9.0	22.0	9.7	31.8	2.3	3.0	3	10.1	0.0	11.4	5.3	9.4	12. 5	6.0	11.3	7.9
DRPC.	Amount	(In wt%)		none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none
TCD***		(In wt%)		none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none
*spuno		Amount	(In wt%)	0.5	0.5	0.5	0.5	0.5	0.5	0.5	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	2.0	2.0	2.0	2.0	2.0	2.0	2.0	3.0	3.0	3.0
Epoxy compounds		Kind		PGE	s-BuPGE	t-BuPGE	DODECYLGE	DECYLGEs	EPCH	EP. DODEC.	PGE	s-BuPGE	t-BuPGE	DODECYLGE	DECYLGES	EPCH	EP. DODEC.	EP. S0Y.	EP. St. B	PGE	s-BuPGE	t-BuPGE	DODECYLGE	DECYLGEs	ЕРСН	EP. DODEC.	PGE	s-BuPGE	t-BuPGE
Base	oils			-	2	e	4	ഹ	9	7	∞	6	10	11	12	13	14	15	16	17	18	19	20	21	22	23		2	
Examples .	Comparative	Examples		Example 1	Example 2	Example 3	Example 4	Example 5	Example 6	Example 7	Example 8	Example 9	Example 10	Example 11	Example 12	Example 13	Example 14	Example 15	Example 16	Example 17	Example 18	Example 19	Example 20	Example 21	Example 22	Example 23	Example 24	Example 25	Example 26

Table 1 (Cont.)

No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change	No change
1	!	-	!	;			i	1		!	1	1	1			;	-	-	-		-		-	1	-	!	}	;	;		-	
	!	:	;	-	;			-	1	-	-	!	1	1 1	† 1	!!!	:	-	-	!	;	-	-	!			-	;	!	1	-	1
1	-	-	-	-	-	-	-		!		ļ	1	1			;		1	-	!	-		:	1	}	i	:	-	-	:	;	-
-		1	-	-	-	-		-		1			-	-	:	-	1	t t	1	1	1	-	:	1 1	!		1	-		1	-	-
0. 18	0.16	0.16	0.12	0.21	0.20	0.22	0.17	0.14	0.15	0.12	0.22	0.21	0.16	0.11	0.17	0.11	0.16	0.20	0.12	0.20	0.18	0.18	0.16	0.16	0.12	0.19	0.19	0.19	0.19	0.19	0.19	0.19
		!	-		-	1	-	;	;	1	:	!	1	ļ	1	1	1	-		-	1	;		!	-	-	•	-	‡ †	:	-	:
2. 9×1014	3.0×1014	3.1×10 ¹⁴	2.4×1014	4.5×1014	4.0×10 ¹⁴	4.1×10 ¹⁴	2.7×10 ¹⁴	2.8×10 ¹⁴	3.1×10 ¹⁴	2.7×10 ¹⁴	4.0×10 ¹⁴	3. 9×10^{14}	2. 4×10^{14}	1.4×1014	2.4×1014	1.1×10 ¹⁴	2.5×10 ¹⁴	2.9×10 ¹⁴	1.4×1014	3.8×10 ¹⁴	3.9×1014	2. 9×10 ¹⁴	3.0×10 ¹⁴	3.1×10 ¹⁴	2.4×1014	4.2×1014	4.2×1014	4.2×1014	4.2×1014	4.2×1014	4.2×1014	4.2×1014
	-	-	;	-	!	1	-	!	!	:	i		-	-	1	-		-	-	1	-	-	:	:	-	;	:	!		-	-	1
	-	1	1	1		;	-	!	1.	i	;	:	1	:	;	i	!	ļ	1	1		:	;	1	-	1	;	1		-	-	1
10.0	28.0	11.6	32. 1	3.5	9.2	6.9	8.8	21.8	9.5	31.6	2.1	2.9	5.3	10.1	6.0	11.4	5.3	9.4	12. 5	11.2	7.9	9.8	28.0	11.6	32.0	0.0	0.0	0.0	0.0	6.0	9.0	6.2
none	попе	none	none	none	none	none	none	none	попе	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none
none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	·none	попе	none	none	none
3.0	3.0	3.0				3.0			3.0				3.0			3.0			3.0				3.0		3.0					3.0	3.0	1.0
DODECYLGE	DECYLGES	ЕРСН	EP. DODEC.	PGE	s-BuPGE	t-BuPGE	DODECYLGE	DECYLGES	ЕРСН	EP. DODEC.	EP. S0Y.	EP. St. B	PGE	s-BuPGE	t-BuPGE	DODECYLGE	DECYLGEs	EPCH	EP. DODEC.	PGE	PGE	PGE	PGE	PGE	PGE	s-BuPGE	t-BuPGE	DODECYLGE	DECYLGES	ЕРСН	EP. DODEC.	PGE
4	ഹ	9	-	<u></u>	6	9	Π	12	13	14	12	16	17	18	19	70	21	77	23	7	ر ى	4	ď	9	7		_	_	_	_		-
Example 27	Example 28	Example 29	Example 30	Example 31	Example 32	Example 33	Example 34	Example 35	Example 36	Example 37	Example 38	Example 39	Example 40	Example 41	Example 42	Example 43	Example 44	Example 45	Example 46	Example 47	Example 48	Example 49	Example 50	Example 51	Example 52	Example 53	Example 54	Example 55	Example 56	Example 57	Example 58	Example 59

Table 1 (Cont.)

Example 60		l PGE	3.0	none	0.2	9.0	1		4. 2×10^{14}	!	0.19			-		No change
Example 61	7	s-BuPGE	3.0	none	0.2	11.3		1	3.8×1014	-	0.20		:	!	1	No change
Example 62	_د	t-BuPGE	3.0	none	0.2	7.9	1	* * *	3, 9×10 ¹⁴	i	0.18	1	1	!		No change
Example 63	4	DODECYLGE	3.0	none	0.2	10.0	;	:	2.9×10 ¹⁴	4 4 1	0.18	-		-	ì	No change
Example 64	ഹ	DECYLGES	3.0	none	0.2	28.0	ł	1	3.0×10 ¹⁴	1	0.16	-	į	-	-	No change
Example 65	9	EPCH	3.0	none	0.2	11.6	;	1 1	3.1×10 ¹⁴	1	0.16	-		:		No change
Example 66	7	EP. DODEC.	3.0	none	0.2	32.0	1	:	2.4×1014	:	0.12			-	1	No change
Example 67	8	PGE	3.0	none	none	5.7	Immiscible -	-24~CT** 2.1×1014	2.1×10 ¹⁴	16	0.21	*	No change	No change	No change	·
Example 68	30	s-BuPGE	3.0	none	none	5.7	Immiscible	-25~CT** 2. 4×1014	2.4×1014	18	0.20		No change	No change	No change	
Example 69	8	t-BuPGE	3.0	none	0.2	5.8	Immiscible	-25~CT** 2. 3×1014	2.3×10 ¹⁴	19	0.21		No change	No change	No change	-

Table 2

Comparative oils -			TCD	DRDC	viscosity	miscibile lemp. Ra	Miscibile Temp. Range mith HFC-1342 (で)	Resisti-	<u></u>	Hygro-			Sealed tube test		
e e	Kind	4 con V	·		0100°C	אַרבוו ווויס דאַ	(C)	025°C	journal	30°C, 60%	011		Catalyst		Appearance
,	MING	(In wt%)	(111 # 0.8)	(III *C\$)	(\$/_ww)	Amount of HFC-134a (90 wt%)	Amount of HFC-134a (97 wt%)	(\$25cm)	worn (mg)	€	color	ŋ	ស	Al	of test oil
Comp. Ex. 1 24 n	none	1	none	none	5.1	Immiscible	1	3.8×10 ¹⁵	23	0.01	1	-			
	none	!	none	none	4.8	Immiscible	!	6. 4×1015	22	0.01	-			-	;
	none	1	none	none	4.9	<-70~97		1.1×10 ¹¹	40	1.31	:	,	1	-	!
	none	-	попе	none	10.8	<-70~56		1.2×10^{11}	32	1.01		1 1	!	:	
	none	!	none	none	10.6	4-70~64		5.6×101°	38	2.30	-	-		1	:
62 59	none	!	none	none	22. 4	-51~32	:	4.8×1010	30	1.81	-	-	-	:	-
	none	:	none	none	6.1	Immiscible	!	2.1×10^{14}	25	0.21	-	No change	Luster decreased	No change	
	none	!	none	none	4.9	Immiscible	:	-	ļ	-	-	1	1	-	
Comp. Ex. 9 32 n	none	1	none	none	5.4	Immiscible	-	-	-	-		† !		-	-
	none	:	none	none	4.4	Immiscible	-	1	-	-	-	: :		-	-
Comp. Ex. 11 1 n	none	1	none	none	6.2	-27~CT**	-45~CT.	4.1×10 ¹⁴	27	0.19	4	No change	Luster decreased	No change	;
12 2	none	!	none	none	11.5	-32~CT**	L2~09-	3.8 $\times 10^{14}$	25	0.19	4	No change	Luster decreased	No change	!
13 3	none	;	none	none	8.1	-28~CT	-45~CT••	4.0×1014	56	0.19	2	No change	Luster decreased	No change	1
14 4	none	1	none	none	10.0	<-70~CT**	<-70~CT**	2. 8×10^{14}	22	0.17	2	No change	Luster decreased	No change	-
15 5	none	:	none	none	28. 2	-1~15	-18~90	3.0×10^{14}	19	0.15	3	No change	Luster decreased	No change	1
16 6	none	:	none	none	11.8	-25~30	-36~CT.	3. 2×10^{14}	22	0.16		No change	Luster decreased	No change	-
17 7	none	;	none	none	32. 3	-5~86	<-70~CT.	2. 4×10 ¹⁴	18	0.13	4	No change	Luster decreased	No change	;
18 8	none	-	none	none	3.8	-35~CT**	<-70~CT**	4.5×10 ¹⁴	29	0.20	4	No change	Luster decreased	No change	:
19 9	none	;	none	none	9.5	-45~CT**	<-70~CT**	3. 9×10^{14}	26	0.20	4	No change	Luster decreased	No change	1 1 1
20 10	none	;	none	none	7.2	-37~CT••	<-70~CT**	4. 1×10^{14}	26	0.20	9	No change	Luster decreased	No change	;
=======================================	none	;	none	none	9.1	<-70~CT.	-25~CT.	2.5×10 ¹⁴	24	0.18	4	No change	Luster decreased	No change	:
22 12	none	1	none	none	22. 1	-11~75	-38~17	2. 8×10^{14}	20	0.14	4	No change	Luster decreased	No change	;
23 13	none	;	none	none	9.8	-20~0	-21~CT**	3.0×10^{14}	24	0.16	2	No change	Luster decreased	No change	;
24 14	none	:	none	none	31.9	-7~88	-19~CT**	2.5×10 ¹⁴	19	0.14		No change	Luster decreased	No change	
25 15	none	-	none	none	2.3	-24~CT.	-38~CT**	4.2×10 ¹⁴	24	0.22	2	No change	Luster decreased	No change	!
Сощр. Ех. 26 16 п	none	i	none	none	3.1	-30~CT**	-54~CT.	3.9×10 ¹⁴	22	0. 20	7	No change	Luster decreased	No change	!

Table 2 (Cont.)

	-			1 1 1			White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid
No change	No change	No change	No change	No change	No change	No change		-	-		-	-		-	-		-	-	-	-	-	-			-	-	-			-		-
Luster decreased	Luster decreased	Luster decreased	Luster decreased	Luster decreased	Luster decreased	Luster decreased		1 1	-		-	-	1	1	-	1	!		1		1 1	1	-	!!!!	!	!	-	f	!!!!	!		-
No change	No change	No change	No change	No change	No change	No change	:	1		!	-	-		1		-	-	-	-		•		-	!	-		1	1	1	:		-
4	4	r	4	2	4			1	-	!	-	-	!	-	-	1	-	-	!	-	1	-	-	-	!	1	-		!	-	1	!
0.18	0.11	0.16	0.10	0.18	0.18	0.12	1.01	1.93	1.93	1.01	1.93	1.01	1.93	1.01	1.93	1.01	1.10	1.07	0.47	0.63	1. 78	0.51	0.74	0.89	09.0	0.59	0.42	0.74	1. 40	1.40	1.40	1.40
15	12	14	10	2	20	17	-	-	-	:	;	!	-	i	-	!		!	:	-	!	-	!	!	!	-	;	;	i	!	!	-
2.5×10¹⁴	1.4×1014	2. 2×1014	1.1×10^{14}	2. 3×10 ¹⁴	2.8×10 ¹⁴	1.2×1014	1. 2×10 ¹¹	6.8×101°	6.8×101°	1.2×10^{11}	6.8×101°	1.2×10^{11}	6.8×101°	1.2×10^{11}	6.8×101°	1.2×10 ¹¹	1.0×10^{11}	1.1×10^{11}	6.8×10 ¹¹	4.8×10 ¹¹	8.4×101°	6. 6×1011	3.8×1011	2. 6×10 ¹¹	3.8×1011	4.1×10 ¹¹	8. 7×10 ¹¹	2.1×10 ¹¹	2.5×10 ¹¹	2.5×10 ¹¹	2.5×10^{11}	2. 5×10 ¹¹
<-70~CT**	-80~CL	<-70~CT.	-56~CT.	<-70~CT	<-50~CT.	<-50~CT.	-			-	-	-	-	-	!	!	-	1					-	-	-	-	-	!	!	-		:
-55~CT.	-34~95	-47~CT.	-32~-92	-55~CT**	<-35~76	<-30~08	-	-	-	-	}		-	-			-	-	-		!	-	!	-			:	;	1		:	!
5.5	10.3	6.2	11.6	5.5	9.6	12.7	10.6	7.0	7.0	10.7	6.9	10.6	6.9	10.7	7.0	10.7	11. 2	12.6	7.2	8.0	6.3	7.6	11.5	11.0	10.4	16.8	11.4	14.4	6.3	6.3	6.3	6.3
none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none
none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none	none
-	}	-	!	-	!	1	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0
none	none	none	none	none	none	none	PGE	PGE	s-BuPGE	t-BuPGE	Podecylge	DECYLGES	EPCH	EP. DODEC.	EP. SOY.	EP. St. B	PGE	PGE	PGE	PGE	PGE	PGE	PGE	PGE	PGE	99.	PGE	PGE	S-BuPGE	t-BuPGE	DODECYLGE	DECYLGEs
17	18	19	50	77	22	23	23	34	34	27	34	27	34	23	34	27	35	98	37	38	33	8	41	42	43	44	45	46	33	39	33	39
Сопр. Ех. 27	Сощр. Ех. 28	Сопр. Ех. 29	Сомр. Ех. 30	Сопр. Ех. 31	Сощр. Ех. 32	Сотр. Ех. 33	Сопр. Ех. 34	Сопр. Ех. 35				Сощр. Ех. 39	Сотр. Ех. 40	Сотр. Ех. 41	Сощр. Ех. 42	Сопр. Ех. 43	Сотр. Ех. 44	Сотр. Ех. 45	Сотр. Ех. 46	Сотр. Ех. 47	Сопр. Ех. 48	Сотр. Ех. 49	Сопр. Ех. 50	Сопр. Ех. 51		Сощр. Ех. 53	Сотр. Ех. 54	Сотр. Ех. 55		Comp. Ex. 57	Сотр. Ех. 58	Сотр. Ех. 59

TOPECT COLUMNIC

2 (Cont.) Tab1e

Г	.g.	<u>.</u>	<u></u>	<u>.</u>	.p	<u>.</u>	<u>.</u>		.9	<u>.</u>	<u></u>
	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid	White-turbid
	:	i	;	ļ	;	:	i	1	1	:	;
	1 1	-	:	-	-	1 1	}	1 4 1	1	!	!
		:		-	1	:	1	-			-
	-				1	1	-	1	1	1	1 1
	1.40	1.40	1.01	1.93	0.47	1.78	0. 22	0.28	0.22	0.28	0.30
	:	-	1 1 1		-	!	}	;	1	1	25
	2.5×10 ¹¹	2.5×10 ¹¹	1.2×10^{11}	6.8×101°	6.8×10 ¹¹	8.4×101°	2.1×10 ¹³	8.6×10 ¹²	2.1×10 ¹³	8.6×10 ¹²	6.2×10 ¹²
	!	;	1	:	!	!	!	!	!	!	-10~54
	i	;	;	-	1	1	1	;	-		Immiscible
	6.3	6.3	10.5	6.9	7.1	6.2	6.1	6.0	6.2	6.1	6.0
	none	none	none	none	попе	none	none	none	none	none	none
	none	none	1.0	1.0	1.0	1.0	1.0	1.0	none	none	none
	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0
	EPCH	EP. DODEC.	PGE	PGE	PGE	PGE	PGE	PGE	PGE	PGE	PGE
_	38	33		34		33			47	48	49
_	Сомр. Ех. 60	Comp. Ex. 61	Сотр. Ех. 62	Comp. Ex. 63	Comp. Ex. 64	Comp. Ex. 65		Comp. Ex. 67	Comp. Ex. 68	Comp. Ex. 69	Сощр. Ех. 70

* PGE : Phenyl Glycidyl Ether

s-BuPGE : s-Butylphenyl Glycidyl Ether

t-BuPGE : t-Butylphenyl Glycidyl Ether

DODECYLGE : Dodecyl Glycidyl Ether

DECYLGEs : Glycidyl 1,1-Dimethyloctanoate

EPCH : 1, 2-, Epoxyclohexane

EP. SOY. : Epoxydized Soy Bean 0il EP. DODEC. : 1, 2-, Epoxydodecane

** CT: Critical temperature of HFC-134a (102°C) EP.St.B : Epoxydized Butyl Stearate

*** TCP: Tricresyl phosphate

**** DBPC; di-tert-butyl-P-cresol

It is apparent from the results indicated in the Tables 1 and 2 that the refrigerator oils (Examples 1-23 and 67-69) of the present invention at least miscibile with 97% by weight of a refrigerant, HFC-134a, and that the refrigerator oils (Examples 1-23) of the present invention are excellent in miscibility with the refrigerant as compared with those of Comparative Examples 1-2.

Examples 3 and 4 as well as the polyoxypropylene glycols of

10 Comparative Examples 5 and 6 are excellent in miscibility
with the refrigerant, but these ethers and glycols are
inferior in insulating property thereby rendering them
unusable for hermetic type compressors. Still further,
Table 2 shows that the ethers and glycols of Comparative

15 Examples 3-6 have hygroscopicity 5-10 times that of the
refrigerator oils of Examples 1-69 and are also inferior in
electrical insulating property, ice choke, wear resistance,
stability and the like to the refrigerator oils of the
Examples.

The FALEX wear test shows that the refrigerator oils of Examples 1-23 and 67-69 are at least equal in wear resistance to those of Comparative Examples 3-6.

The refrigerator oils of Comparative Examples 7 and 11-33 which are free from epoxy compounds are equivalent to those of Examples 1-23 and 67-69 of the present invention in miscibility with HFC-134a, insulating property, wear resistance and hygroscopicity, but, as is apparent from the

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results of the sealed tube test, the refrigerator oils of Comparative Examples 7 and 11-33 are quite inferior in thermal and chemical stability as compared with the refrigerator oils of the present invention.

each of which contains at least a polypropylene glycol monoalkyl ether or a derivative thereof and an epoxy compound have a disadvantage that a precipitate which is insoluble into the base oil and the refrigerant is formed in the refrigerator oils due to deterioration of the epoxy compound during the storage under a high humidity. The formed precipitate makes the refrigerator oil white-turbid and leads to danger of blocking an expansion valve and/or a capillary tube in a refrigerator system in which the above refrigerator oils of Comparative Examples 34-70 is used. Further, the refrigerator oils of Comparative Examples 34-70 have lower resistivity and higher hygroscopicity than any one of refrigerator oils of the present invention.

On the other hand, the refrigerator oils (Examples 20 24-69) of the present invention are free from the generation of such a precipitate even if they are stored under a high humidity. Therefore, in a case where the refrigerator oil of the present invention is used in a refrigerator system, there is no danger as mentioned above.

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(Effect of the Invention)

As is apparent from the above comparative experiments, the refrigerator oils of the present invention are suitable for use in refrigerators using therein a hydrogen-containing halogenocarbon as a refrigerant and are excellent in electrical insulating property, wear resistance, nonhygroscopicity and thermal and chemical stability. In other words, the refrigerator oils of the present invention excellently have all properties which are important for refrigerator oils.